

UNITED STATES DISTRICT COURT
DISTRICT OF NEW JERSEY

CHAMBERS OF
ESTHER SALAS
UNITED STATES DISTRICT JUDGE

MARTIN LUTHER KING
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December 13, 2017

LETTER ORDER re: ITINERARY FOR ORAL ARGUMENT

**Re: *Mylan Pharmaceuticals Inc. v. Celgene Corporation*,
Civil Action No. 14-2094 (ES) (MAH)**

Dear Counsel:

The Court anticipates that oral argument will proceed in the following order:

- (1) Does the “rule of reason” burden-shifting framework apply? If so, how does that framework apply in the context of the pending summary judgment motion?
- (2)
 - a. Can Celgene be held liable for conduct before FDA “approval” of Mylan’s protocols?
 - b. Can Celgene be held liable for conduct after FDA “approval” of Mylan’s protocols?
- (3)
 - a. What is Mylan’s current damages theory for Revlimid®? In particular, what does Exhibit 99 provide in that regard?
 - b. What is Mylan’s new request for a damages theory relating to Revlimid®—and why should the Court consider it?
- (4) Does Celgene’s statute-of-limitations defense bar Thalomid®-related claims? In particular, is there a reason based in law or evidence supporting that “damages for generic entry are not ‘unascertainable’ up to the moment of market entry”? (See D.E. No. 240 at 19).
- (5) Is there a genuine issue of material fact that Mylan would have “skinny labeled” in light of Celgene’s orphan drug exclusivity for Thalomid®?

- (6) If the Court finds that Celgene *can* be held liable for conduct *after* FDA “approval” of Mylan’s protocols, do Mylan’s New Jersey law claims necessarily survive?

SO ORDERED.

s/Esther Salas
Esther Salas, U.S.D.J.